K024130

SPECIAL 510(k) SUMMARY

DEC 3 0 2002

1.0 Submitter:

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Date of Summary Prepared: 13 DEC 2002

2.0 Contact Person:

Name: Mr. Terence Lim
Phone No.: +60 3 8706 1486
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3.0 Modified Device Identification:

Trade Name: 1) Comfit, and
2) Multiple or Customers' Trade Name
Device Name: Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile
Common Name: Surgical Gloves
Classification Name: Surgeon's Gloves (per 21 CFR 878.4460)

4.0 Identification of the Legally Marketed Device:

Class I Powder Free synthetic rubber latex surgeon's gloves, 79KGO, that meets all the requirements of ASTM standard D 3577 – 01a^{E2} Type 2 and FDA 21 CFR 800.20.

5.0 Description of Device Modification:

The Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile is equivalent to the existing model, i.e. Profeel Powder Free Polymer Coated Neoprene Surgical Gloves, Sterile which had submitted and cleared under 510(k) number K003019.

The difference in this submission is:

a) Change of colour additive used, i.e. Green

The modification of colour does not affect the intended use of the device as well as it does not affect its safety and effectiveness. The indication for use and proposed labeling for the device are illustrated in subsequent sections.

The Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile meets all the requirements of ASTM standard D 3577 – 01a^{E2} and FDA 21 CFR 800.20.

6.0 Intended Use of the Device:

The Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile are made of synthetic rubber latex intended to be worn on the hand of healthcare personnel, operating room personnel and similar personnel to prevent contamination between the healthcare or similar personnel and the patient's body, fluids, waste, or environment.

7.0 Summary of Technological Characteristics for the Modified Device:

The Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile are summarized with the following technological characteristics compared to ASTM or equivalent standards.

CHARACTERISTICS	STANDARDS	DEVICE PERFORMANCE
Dimensions	ASTM D 3577 – 01a^{E2}	Meets
Physical Properties	ASTM D 3577 – 01a^{E2}	Meets
Freedom from pinholes	ASTM D 3577 – 01a^{E2} FDA 21 CFR 800.20	Meets
Powder Residual	ASTM D 6124 – 01	Meets < 2 mg/glove

8.0 Conclusion:

The Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile will perform according to the glove performance standards referenced in section 7 above and meet ASTM standards, and FDA requirements for waterleak test on pinhole AQL. Consequently, this device is substantially equivalent to currently marketed devices.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

DEC 3 0 2002

Mr. Terence Lim
Associate Manager, RA/QA
WRP Asia Pacific Sdn Bhd
Lot 1, Jalan 3, Kawasan Perusahaan
Bandar Baru Salak Tinggi,
43900 Sepang,
Selangor Darul Ehsan,
MALAYSIA

Re: K024130
Trade/Device Name: Powder Free Polymer Coated Green Neoprene Surgical Gloves, Sterile
Regulation Number: 21 CFR 878.4460
Regulation Name: Surgical Gloves
Regulatory Class: I
Product Code: KGO
Dated: December 13, 2002
Received: December 16, 2002

Dear Mr. Lim:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4618. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address http://www.fda.gov/cdrh/dsma/dsmamain.html

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Anesthesiology, General Hospital,
Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

WRP Asia Pacific Sdn Bhd

147817V

INDICATIONS FOR USE

Applicant: WRP Asia Pacific Sdn Bhd

510(k) Number (if known): K024130

Device Name: POWDER FREE POLYMER COATED GREEN NEOPRENE SURGICAL GLOVES, STERILE

Indications For Use:

The surgical glove is a device made of synthetic latex intended to be worn by surgeons and/or operating room personnel to protect a surgical wound from contamination.

_____ _____ _____ _____ _____

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use _____ OR Over-The-Counter _____
(Per 21 CFR 801.109)

Chin S. Lin

(Division Sign-Off)
Division of Anesthesiology, General Hospital, Infection Control, Dental Devices

510(k) Number K024130

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